



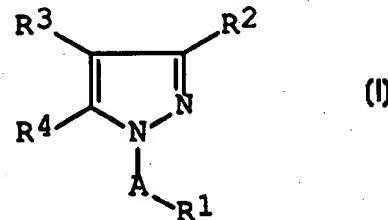
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(54) Title: 5-ARYLPYRAZOLE COMPOUNDS

(57) Abstract

A compound of formula (I), wherein R¹ is aryl optionally substituted with substituent(s) selected from the group consisting of halogen, nitro, amino, esterified carboxy, carboxy, carbamoyl, cyano and lower alkoxy, R² is hydrogen, amino, halogen, esterified carboxy, carboxy, carbamoyl, cyano, or lower alkyl substituted with halogen, R³ is hydrogen, aryl optionally substituted with halogen, or lower alkyl optionally substituted with hydroxy, amino or carboxy, R⁴ is aryl substituted with substituent(s) selected from the group consisting of halogen, lower alkoxy, lower alkylthio, lower alkylsulfinyl, lower alkylsulfonyl, sulfamoyl, and lower alkylsulfamoyl, and A is lower alkylene, and its salt, which are useful as medicament.



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DESCRIPTION

5-ARYLPYRAZOLE COMPOUNDS5 Technical Field

This invention relates to 5-arylpyrazole compounds having pharmacological activity, to a process for their production and to a pharmaceutical composition containing the same.

10

Background Art

Some pyrazole derivatives having antiinflammatory and/or analgesic activities have been known, for example, in Canadian Patent 1 130 808, and EP Patent Publication Nos. 248 15 594, 272 704, 293 220, 418 845 and 554 829, and WO Patent Publication Nos. 95/15315, 95/15316, 95/15317, 95/15318, 96/14302 and 97/15271.

Disclosure of Invention

20 This invention relates to 5-arylpyrazole compounds, which have pharmaceutical activity such as cyclooxygenase-2 (hereinafter described as COX-II) inhibiting activity, to a process for their production, to a pharmaceutical composition containing the same and to a use thereof.

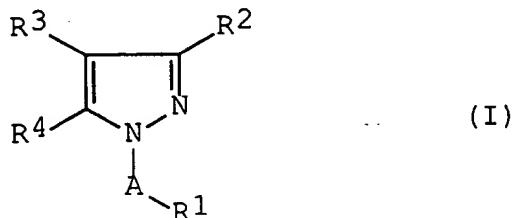
25 Accordingly, one object of this invention is to provide the 5-arylpyrazole compounds, which have a COX-II inhibiting activity.

Another object of this invention is to provide a process for production of the 5-arylpyrazole compounds.

30 A further object of this invention is to provide a pharmaceutical composition containing, as active ingredients, the 5-arylpyrazole compounds.

Still further object of this invention is to provide a use of the 5-arylpyrazole compounds for manufacturing a 35 medicament for treating or preventing various diseases.

The new 5-arylpyrazole compounds of this invention can be represented by the following general formula (I):



wherein R¹ is aryl optionally substituted with substituent(s) selected from the group consisting of halogen, 10 nitro, amino, esterified carboxy, carboxy, carbamoyl, cyano and lower alkoxy,

R² is hydrogen, amino, halogen, esterified carboxy, carboxy, carbamoyl, cyano, or lower alkyl substituted with halogen,

15 R³ is hydrogen, aryl optionally substituted with halogen, or lower alkyl optionally substituted with hydroxy, amino or carboxy,

R⁴ is aryl substituted with substituent(s) selected 20 from the group consisting of halogen, lower alkoxy, lower alkylthio, lower alkylsulfinyl, lower alkylsulfonyl, sulfamoyl, and lower alkylsulfamoyl, and

A is lower alkylene,

25 and its salt.

The compounds of formula (I) may contain one or more asymmetric centers and thus they can exist as enantiomers or diastereoisomers. This invention includes both mixtures and 30 separate individual isomers.

The compounds of the formula (I) may also exist in tautomeric forms and the invention includes both mixtures and separate individual tautomers.

The compound of the formula (I) and its salt can be in a 35 form of a solvate, which is included within the scope of the

present invention. The solvate preferably include a hydrate and an ethanolate.

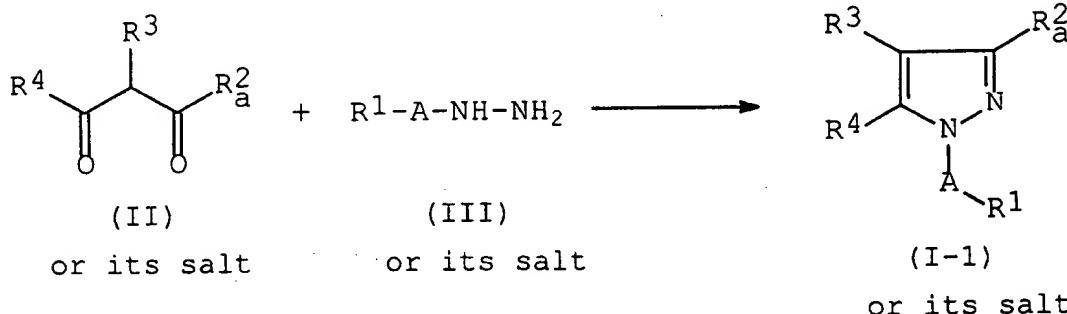
Also included in the scope of invention are radiolabelled derivatives of compounds of formula (I) which 5 are suitable for biological studies.

The object compound (I) or its salt can be prepared by the following process.

10

Process 1

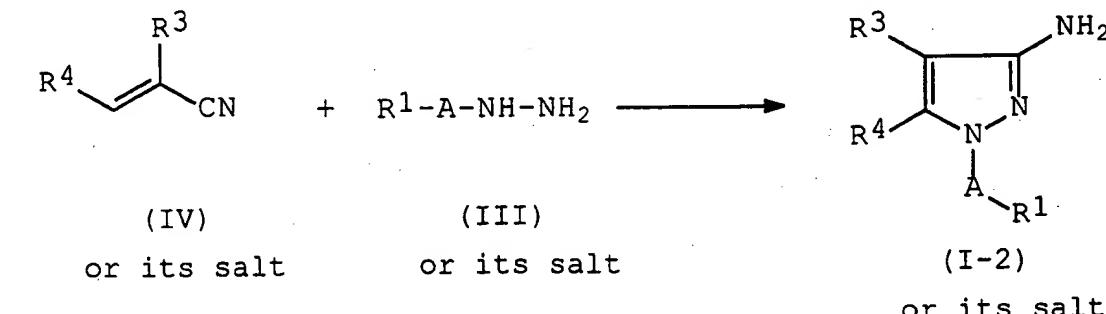
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Process 2

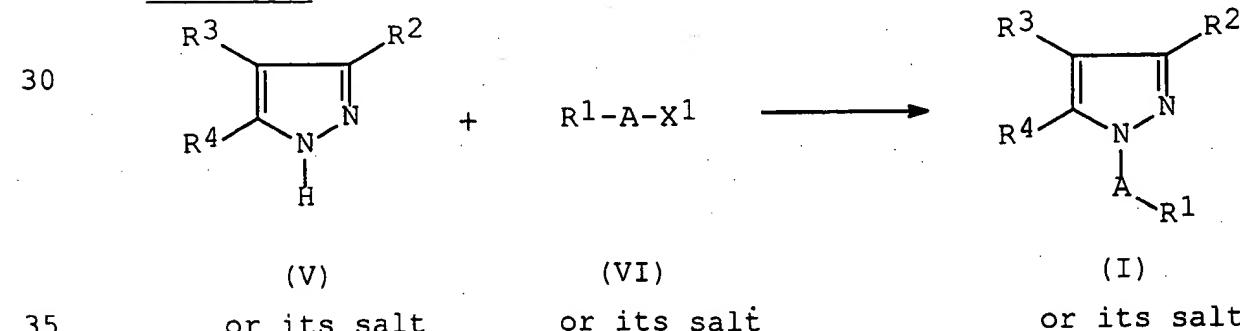
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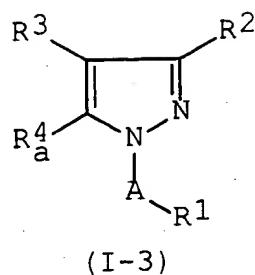
Process 3

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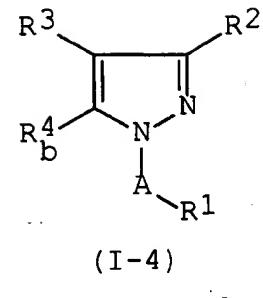


Process 4

5



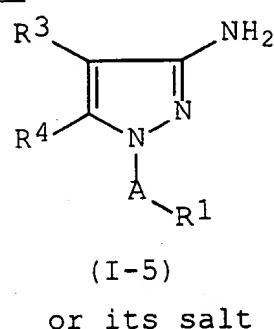
Oxidation



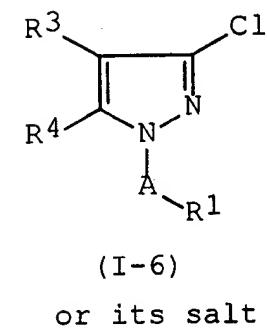
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Process 5

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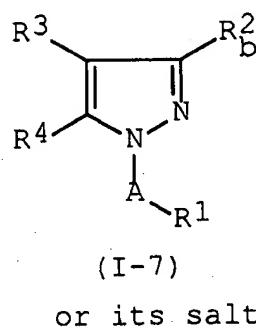
Chlorination



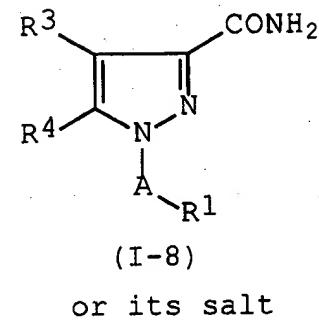
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Process 6

25



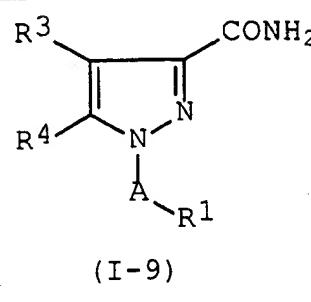
Amidation



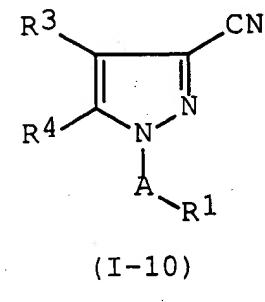
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Process 7

35



Amidation



wherein R^1 , R^2 , R^3 , R^4 and A are each as defined above,

R_a^2 is hydrogen, esterified carboxy, carbamoyl, cyano, or lower alkyl substituted with halogen,

5 R_b^2 is esterified carboxy or carboxy,

R_a^4 is aryl substituted with lower alkylthio,

R_b^4 is aryl substituted with lower alkylsulfinyl or lower alkylsulfonyl, and

x^1 is a leaving group.

10

In the above and subsequent description of the present specification, suitable examples of the various definitions to be included within the scope of the invention are explained in detail in the following.

15

The term "lower" is intended to mean a group having 1 to 6 carbon atom(s), unless otherwise provided.

Suitable "lower alkyl" and lower alkyl moiety in the terms "lower alkylthio", "lower alkylsulfinyl", "lower alkylsulfonyl", "lower alkylsulfamoyl" and "lower alkoxy" may 20 be a straight or branched one, such as methyl, ethyl, propyl, isopropyl, butyl, isobutyl, tert-butyl, pentyl, hexyl, or the like, in which preferable one is methyl.

Suitable "lower alkyl substituted with halogen" may be 25 difluoromethyl, trifluoromethyl, or the like.

Suitable "lower alkylene" may be straight or branched one having 1 to 6 carbon atom(s), such as methylene, ethylene, trimethylene, hexamethylene, or the like, preferably one having 1 to 3 carbon atom(s), more preferably 30 methylene.

Suitable "halogen" may be fluoro, chloro, bromo or iodo.

Suitable "aryl" may be phenyl, naphtyl, tolyl, xylyl, ethylphenyl, propylphenyl, or the like.

Suitable "esterified carboxy" may be substituted or 35 unsubstituted lower alkoxy carbonyl [e.g., methoxycarbonyl,

ethoxycarbonyl, propoxycarbonyl, butoxycarbonyl, 2,2,2-trichloroethoxycarbonyl, etc.], substituted or unsubstituted aryloxycarbonyl [e.g., phenoxy carbonyl, 4-nitrophenoxy carbonyl, etc.], substituted or unsubstituted 5 ar(lower)alkoxycarbonyl [e.g. benzyloxycarbonyl, 4-nitrobenzyloxycarbonyl, etc.], or the like.

Suitable "leaving group" may be halogen, acyloxy [e.g., acetyloxy, trifluoroacetyloxy, etc.], lower alkylsulfonyloxy [e.g., methanesulfonyloxy, etc.], triarylphosphinoxy [e.g., 10 $-O-P^+(C_6H_5)_3$, etc.], or the like.

Suitable salts of the compounds (I) and (I-1) to (I-10), and the compounds (II) to (VI) are pharmaceutically acceptable conventional non-toxic salts and include a metal 15 salt such as an alkali metal salt (e.g., sodium salt, potassium salt, etc.) and an alkaline earth metal salt (e.g., calcium salt, magnesium salt, etc.), an ammonium salt, an organic base salt (e.g., trimethylamine salt, triethylamine salt, pyridine salt, picoline salt, dicyclohexylamine salt, 20 etc.), an organic acid salt (e.g., acetate, maleate, tartrate, methanesulfonate, benzenesulfonate, formate, toluenesulfonate, trifluoroacetate, etc.), an inorganic acid salt (e.g., hydrochloride, hydrobromide, sulfate, phosphate, etc.), a salt with an amino acid (e.g., arginine, aspartic 25 acid, glutamic acid, etc.), or the like.

The processes for preparing the object compound (I) are explained in detail in the following.

30 Process 1

The compound (I-1) or its salt can be prepared by reacting the compound (II) or its salt with a hydrazine derivative (III) or its salt.

35 The reaction is usually carried out in a conventional solvent such as water, alcohol [e.g., methanol, ethanol,

isopropyl alcohol, etc.], alkanoic acid, tetrahydrofuran, dioxane, toluene, methylene chloride, chloroform, N,N-dimethylformamide, or any other organic solvents which do not adversely affect the reaction, or the mixture thereof, 5 preferably, acidic solvent such as alkanoic acid (e.g., acetic acid).

The reaction temperature is not critical and the reaction is usually carried out under cooling to warming.

10 Process 2

The compound (I-2) or its salt can be prepared by reacting the compound (IV) or its salt with the compound (III) or its salt.

15 This reaction can be carried out in a similar manner to that of Process 1, and therefore the reaction conditions (e.g., solvent, reaction temperature, etc.) can be referred to those of the Process 1.

20 Process 3

The compound (I) or its salt can be prepared by reacting the compound (V) or its salt with the compound (VI) or its salt.

This reaction is usually carried out in the presence of an inorganic or an organic base.

25 Suitable inorganic base may include an alkali metal [e.g., sodium, potassium, etc.], an alkali metal hydroxide [e.g., sodium hydroxide, potassium hydroxide, etc.], alkali metal hydrogen carbonate [e.g., sodium hydrogen carbonate, potassium hydrogen carbonate, etc.], alkali metal carbonate [e.g., sodium carbonate, etc.], alkali earth metal carbonate [calcium carbonate, etc.], or the like.

30 Suitable organic base may include tri(lower)alkylamine [e.g. triethylamine, N,N-diisopropylethylamine, etc.], alkyl lithium [e.g. methyl lithium, butyl lithium, etc.], lithium diisopropylamide, lithium hexamethyldisirazido, alkali metal

hydride [e.g., sodium hydride, potassium hydride, etc.] or the like.

The reaction is usually carried out in a conventional solvent such as water, alcohol [e.g., methanol, ethanol, 5 isopropyl alcohol, etc.], tetrahydrofuran, dioxane, toluene, methylene chloride, chloroform, N,N-dimethylformamide or any other organic solvents which do not adversely affect the reaction, or the mixture thereof.

10 The reaction temperature is not critical and the reaction is usually carried out under cooling to warming.

Process 4

15 The compound (I-4) or its salt can be prepared by reacting a compound (I-3) or its salt with an oxidizing agent.

The suitable oxidizing agent may be hydrogen peroxide, cumene hydroperoxide, tert-butyl hydroperoxide, Jones reagent, peracid [e.g. peracetic acid, perbenzoic acid, m-chloroperbenzoic acid, monopersulfate compound 20 (oxone[®]), etc.], chromic acid, potassium permanganate, alkali metal periodate [e.g. sodium periodate, etc.], and the like.

25 This reaction is usually carried out in a solvent which does not adversely influence the reaction such as acetic acid, dichloromethane, acetone, ethyl acetate, chloroform, water, an alcohol [e.g. methanol, ethanol, etc.], a mixture thereof or the like.

The reaction temperature is not critical, and the reaction is usually carried out under cooling to warming.

30 Process 5

The compound (I-6) or its salt can be prepared from the compound (I-5) or its salt by the following methods.

Namely, 1) the compound (I-5) or its salt is firstly reacted with a nitrite compound, and then 2) the resulting 35 product is reacted with cuprous chloride.

Suitable nitrite compound may be alkali metal nitrite [e.g. sodium nitrite, potassium nitrite, etc.], alkyl nitrite [e.g. isoamyl nitrate, tert-butyl nitrite, etc.], and the like.

5 In the first step, the reaction is preferably carried out in the presence of an acid [e.g. hydrochloric acid sulfuric acid, etc.].

10 The reaction is usually carried out in a solvent such as water, tetrahydrofuran, dioxane, acetonitrile, or any other organic solvent which does not adversely influence the reaction, or a mixture thereof.

The reaction temperature is not critical and the reaction can be carried out under cooling to warming.

15 In the second step, the reaction is preferably carried out in the presence of alkali metal halide [e.g. sodium chloride, etc.] and an inorganic acid [e.g. hydrochloric acid, etc.].

20 The reaction is usually carried out in a solvent such as water, tetrahydrofuran, dioxane, or any other organic solvent which does not adversely influence the reaction, or a mixture thereof.

The reaction temperature is not critical and the reaction can be carried out warming to heating.

25 Process 6

The compound (I-8) or its salt can be prepared by subjecting the compound (I-7) or its salt to amidation reaction.

30 Amidation reaction can be carried out in a conventional manner, which is capable of converting carboxy group or protected carboxy group to carbamoyl group. Amidation can preferably carried out by, for example, (i) reacting the compound (I-7), wherein R_b^2 is esterified carboxy, with alkanoylamine (e.g., aceamide, formamide, etc.) in the presence of organic base (e.g., sodium alkoxide, etc.) or

(ii) reacting the compound (I-7), wherein R_b^2 is carboxy, or its salt, with ammonia or its salt in the presence of condensing agent.

The reaction is usually carried out in a conventional solvent such as alcohol [e.g., methanol, ethanol, isopropyl alcohol, etc.], tetrahydrofuran, dioxane, toluene, methylene chloride, chloroform, N,N-dimethylformamide, or any other organic solvents which do not adversely affect the reaction, or the mixture thereof.

10 The reaction temperature is not critical and the reaction is usually carried out under cooling to warming.

Process 7

15 The compound (I-10) or its salt can be prepared by subjecting the compound (I-9) or its salt to dehydration reaction.

20 Dehydration reaction can be carried out in the conventional manner, which is capable of dehydrating a carbamoyl group to cyano group, and suitable dehydrating agent may be phosphorus compound (e.g., phosphorus oxychloride, etc.) or the like.

25 The reaction is usually carried out in a conventional solvent such as alcohol [e.g., methanol, ethanol, isopropyl alcohol, etc.], tetrahydrofuran, dioxane, toluene, methylene chloride, chloroform, N,N-dimethylformamide, or any other organic solvents which do not adversely affect the reaction, or the mixture thereof.

30 The reaction temperature is not critical and the reaction is usually carried out under cooling to warming.

35

Suitable organic base may be tri(lower)alkylamine [e.g., triethylamine, N,N-diisopropylethylamine, etc.], alkyl magnesium bromide [e.g., methyl magnesium bromide, ethyl magnesium bromide, etc.], alkyl lithium [e.g., methyl lithium, butyl lithium, etc.], lithium diisopropylamide,

lithium hexamethyldisirazido, or the like.

The reaction is usually carried out in a conventional solvent such as water, alcohol [e.g., methanol, ethanol, isopropyl alcohol, etc.], tetrahydrofuran, dioxane, toluene, 5 methylene chloride, chloroform, N,N-dimethylformamide, or any other organic solvents which do not adversely affect the reaction, or the mixture thereof.

The reaction temperature is not critical and the reaction is usually carried out under cooling to warming.

10

The object compound (I) or pharmaceutically acceptable salts thereof of this invention possesses COX-II inhibiting activity and possesses strong antiinflammatory, antipyretic, analgesic, antithrombotic, anti-cancer activities, and so on.

15 The object compound (I) and pharmaceutically acceptable salt thereof, therefore, are useful for treating and/or preventing COX-II mediated diseases, inflammatory conditions, various pains, collagen diseases, autoimmune diseases, various immunological diseases, thrombosis, cancer and 20 neurodegenerative diseases in human beings or animals by using administered systemically or topically. More particularly, the object compound (I) and pharmaceutically acceptable salts thereof are useful for treating and/or preventing inflammation and pain in joint and muscle [e.g. 25 rheumatoid arthritis, rheumatoid spondylitis, osteoarthritis, gouty arthritis, juvenile arthritis, etc.], inflammatory skin condition [e.g. sunburn, burns, eczema, dermatitis, etc.], inflammatory eye condition [e.g. conjunctivitis, etc.], lung disorder in which inflammation is involved [e.g. asthma, 30 bronchitis, pigeon fancier's disease, farmer's lung, etc.], condition of the gastrointestinal tract associated with inflammation [e.g. aphthous ulcer, Crohn's disease, atopic gastritis, gastritis varialoforme, ulcerative colitis, coeliac disease, regional ileitis, irritable bowel syndrome, 35 etc.], gingivitis, inflammation, pain and tumescence after

operation or injury, pyrexia, pain and other conditions associated with inflammation, particularly those in which lipoxygenase and cyclooxygenase products are a factor, systemic lupus erythematosus, scleroderma, polymyositis, 5 tendinitis, bursitis, periarthritis nodose, rheumatic fever, Sjögren's syndrome, Behcet disease, thyroiditis, type I diabetes, nephrotic syndrome, aplastic anemia, myasthenia gravis, uveitis contact dermatitis, psoriasis, Kawasaki disease, sarcoidosis, Hodgkin's disease, Alzheimers disease, 10 or the like. Additionally, the object compound (I) or a salt thereof is expected to be useful as therapeutical and/or preventive agents for cardiovascular or cerebrovascular diseases, the diseases caused by hyperglycemia and hyperlipemia.

15

The patents, patent applications and publications cited herein are incorporated by reference.

20 In order to illustrate the usefulness of the object compound (I), the pharmacological test data of the compound (I) are shown in the following.

[A] ANTIINFLAMMATORY ACTIVITY :

25

Effect on adjuvant arthritis in rats :

(i) Test Method :

30 Ten female Sprague-Dawley rats were used per group. A dose of 0.5 mg of *Mycobacterium tuberculosis* (strain M37 BA) suspended in 0.05 ml of liquid paraffin was injected subcutaneously in the right hind paw. The injection of mycobacterial adjuvant produced local inflammatory lesions (primary lesion) and then about 10 days later, secondary 35 lesions in both the injected and uninjected paws. The

volumes of both paws before and on days 23 after the injection was measured as percent inhibition in comparison to vehicle-treated controls. The drug was given orally once a day for 23 consecutive days from day 1 after the injection.

5

(ii) Test Results :

Test compound (Example No.)	Dose (mg/kg)	Inhibition of secondary lesion (uninjected paw) (%)
10	3	>60

[B] COX-I and COX-II activity in vitro :

(i) Test Method :

15

Cyclooxygenase activities in the absence or presence of inhibitors were measured by determining the level of prostaglandin E₂ (PGE₂) synthesis from arachidonic acid. Enzymes (1 µg for COX-I and/or 3 µg for COX-II) in a total 20 volume of 200 µl of reaction buffer were incubated in the absence or presence of various concentrations of inhibitors for 5 minutes at 30°C. The reaction was then started by the addition of arachidonic acid to the final concentration of 10 µM. The reaction was terminated by 50 µl of 1N-HCl after 25 incubation at 30°C for 5 minutes. PGE₂ was extracted with ethyl acetate, concentrated under a stream of nitrogen and analyzed by a radio immunoassay kit (Amersham) according to the manufacturer's instructions.

COX inhibiting activity of the test compound was assayed 30 as PGE₂ formation using radioimmunoassay to detect a prostaglandin release. The appropriate COX enzyme was incubated in 0.1 M Tris-HCl buffer (pH 7.3) containing hematin and tryptophan with the addition of arachidonic acid (10 µM) for 5 minutes at 37°C. Compounds were pre-incubated

with the enzyme for 5 minutes prior to the addition of arachidonic acid. Any reaction between the arachidonic acid and the enzyme was stopped after 5 minutes at 37°C by addition of 20 μ l of 1N HCl. PGE₂ formation was measured by 5 radioimmunoassay (Amersham).

It appeared, from the below-mentioned Test Results, that the compound (I) or pharmaceutically acceptable salts thereof of the present invention have a selective inhibiting activity against COX-II.

10

(ii) Test Results :

Test Compound (Example No.)	Human COX-I IC ₅₀ (μ M)	Sheep COX-II IC ₅₀ (μ M)
15 4-(2)	>100	< 1
5-(4)	> 15	< 1

For therapeutic purpose, the compound (I) and a pharmaceutically acceptable salt thereof of the present 20 invention can be used in a form of pharmaceutical preparation containing one of said compounds as an active ingredient, in admixture with a pharmaceutically acceptable carrier such as an organic or inorganic solid or liquid excipient suitable for oral, parenteral or external administration. The 25 pharmaceutical preparations may be capsules, tablets, dragees, granules, inhalant, suppositories, solution, lotion, suspension, emulsion, ointment, gel, or the like. If desired, there may be included in these preparations, auxiliary substances, stabilizing agents, wetting or 30 emulsifying agents, buffers and other commonly used additives.

While the dosage of therapeutically effective amount of the compound (I) will vary depending upon the age and

condition of each individual patient, an average single dose of about 0.01 mg, 0.1 mg, 1 mg, 10 mg, 50 mg, 100 mg, 250 mg, 500 mg and 1000 mg of the compound (I) may be effective for treating the above-mentioned diseases. In general, amounts 5 between 0.01 mg/body and about 1,000 mg/body may be administered per day.

The following Preparation and Examples are given for the purpose of illustrating the present invention in detail.

Preparation 1

Sodium hydride (60 %, 1.34 g) was added to a solution of 4-acetylbenzenesulfonamide (1.99 g) and ethyl trifluoroacetate (3.26 g) in dimethylformamide (20 ml) at 5 10-15°C. The resultant mixture was stirred at room temperature for 1 hour and poured into a mixture of ethyl acetate and 1N-hydrogen chloride solution. The organic layer was washed with saturated sodium hydrogen carbonate, and brine successively, dried, and concentrated to dryness. 10 The residue was triturated with isopropyl ether to afford 4-(4,4,4-trifluoro-3-oxo-butyryl)benzenesulfonamide (0.95 g) as a pale yellow crystals.

mp : 142-144 °C

IR (KBr) : 1778, 1602, 1565, 1457 cm^{-1}

15 NMR (DMSO-d₆, δ) : 2.52 (3H, s), 7.03 (1H, s), 7.50- 8.02 (7H, m)

Preparation 2

A mixture of 3',4'-difluoroacetophenone (23.8 g) and 20 sodium thiomethoxide (95 %, 12.4 g) in acetonitrile (720 ml) was stirred at room temperature for 4 hours and warmed at 50°C for 2 hours. The reaction mixture was poured into a mixture of ethyl acetate and water. The organic layer was washed with water, and brine successively, dried, and 25 concentrated to dryness. The residue was triturated with n-hexane to afford 3'-fluoro-4'-(methylthio)acetophenone (17.5 g) as a pale yellow crystals.

mp : 70-72 °C

IR (KBr) : 1673, 1596, 1558, 1403 cm^{-1}

30 NMR (DMSO-d₆, δ) : 2.52 (3H, s), 3.33 (3H, s), 7.50- 7.90 (3H, m)

Preparation 3

35 The following compound was prepared in a similar manner to that of Preparation 1.

4,4,4-Trifluoro-1-[3-fluoro-4-(methylthio)phenyl]-butane-1,3-dione

pale yellow crystals

mp : 60-63 °C

5 IR (KBr) : 1641, 1602, 1548, 1464 cm⁻¹

NMR (CDCl₃, δ) : 2.54 (3H, s), 6.51 (1H, s), 7.50-7.90 (3H, m)

Preparation 4

10 The following compound was prepared in a similar manner to that of Example 2.

4,4,4-Trifluoro-1-[3-fluoro-4-(methylsulfonyl)-phenyl]butane-1,3-dione

pale yellow crystals

15 mp : 109-110 °C

IR (KBr) : 1689, 1683, 1571, 1411 cm⁻¹

NMR (CDCl₃, δ) : 3.27 (3H, s), 6.58 (1H, s), 7.80-8.20 (3H, m)

Example 1

20 A mixture of 4,4,4-trifluoro-1-[4-(methylthio)-phenyl]butane-1,3-dione (405 mg) and 4-chlorobenzyl-hydrazine (300 mg) in acetic acid (5 ml) was refluxed for 1 hour. The solvent was evaporated under reduced pressure and ethyl acetate and water were added to the residue. The organic layer was washed with saturated sodium hydrogen carbonate, and brine successively, dried, and concentrated to dryness. The residue was chromatographed (toluene) over silica gel to afford 1-(4-chlorobenzyl)-5-[4-(methylthio)-phenyl]-3-(trifluoromethyl)pyrazole (0.35 g) as a pale yellow oil.

IR (neat) : 1621, 1590, 1490 cm⁻¹

NMR (CDCl₃, δ) : 2.51 (3H, s), 3.80 (2H, s), 6.06 (1H, s), 7.00-8.00 (8H, m)

35 MS : 383 [M+H]⁺

Example 2

m-Chloroperbenzoic acid (80%, 360 mg) was added to a solution of 1-(4-chlorobenzyl)-5-[4-(methylthio)phenyl]-3-(trifluoromethyl)pyrazole (260 mg) in methylene chloride (20 ml) at 0-5°C. The reaction mixture was stirred for 1 hour, and then washed with saturated sodium hydrogen carbonate, and brine successively, and dried. After removal of the solvent, the residue was chromatographed (toluene : ethyl acetate = 5 : 1) over silica gel to afford 1-(4-chlorobenzyl)-5-[4-(methylsulfonyl)phenyl]-3-(trifluoromethyl)pyrazole (60 mg) as a white crystals.

mp : 112-115 °C

IR (KBr) : 1693, 1496, 1409 cm⁻¹

15 NMR (CDCl₃, δ) : 3.09 (3H, s), 5.34 (2H, s), 6.67 (1H, s), 6.80-7.40 (4H, m), 7.36 (2H, d, J=8Hz), 8.02 (2H, d, J=8Hz)

MS : 415(M⁺), 417(M⁺²)

20 Example 3

A mixture of 4,4,4-trifluoro-1-[4-(methylsulfonyl)phenyl]butane-1,3-dione (1.3 g) and 3-fluorobenzylhydrazine (774 mg) in acetic acid (15 ml) was refluxed for 2 hours.

25 The solvent was evaporated under reduced pressure, and ethyl acetate and water were added to the residue. The organic layer was washed with saturated sodium hydrogen carbonate, and brine successively, dried, and concentrated to dryness. The residue was recrystallized from ethanol to afford 1-(3-fluorobenzyl)-5-[4-(methylsulfonyl)phenyl]-3-(trifluoromethyl)pyrazole (0.76 g) as a white crystals.

mp : 120-122 °C

IR (KBr) : 1594, 1496, 1479 cm⁻¹

35 NMR (CDCl₃, δ) : 3.10 (3H, s), 5.37 (2H, s), 6.69 (1H, s), 6.70-7.20 (4H, m), 7.49 (2H, d, J=8Hz), 8.00 (2H, d, J=8Hz)

MS : 399 [M+H]⁺

Example 4

5 The following compounds described in (1)-(2) were prepared in a similar manner to that of Example 1.

(1) 1-(4-Fluorobenzyl)-5-[4-(methylsulfonyl)phenyl]-3-(trifluoromethyl)pyrazole
white crystals
10 mp : 140-142 °C
IR (KBr) : 1608, 1509, 1467 cm⁻¹
NMR (CDCl₃, δ) : 3.10 (3H, s), 5.34 (2H, s), 6.69 (1H, s), 6.90-7.30 (4H, m), 7.49 (2H, d, J=8Hz), 8.01 (2H, d, J=8Hz)
15 MS : 399 [M+H]⁺

(2) 1-Benzyl-5-[4-(methylsulfonyl)phenyl]-3-(trifluoromethyl)pyrazole
white crystals.
20 mp : 123-125 °C
IR (KBr) : 1606, 1502, 1463 cm⁻¹
NMR (CDCl₃, δ) : 3.08 (3H, s), 5.39 (2H, s), 6.68 (1H, s), 6.90-7.40 (4H, m), 7.48 (2H, d, J=8Hz), 7.98 (2H, d, J=8Hz)
25 MS : 381 [M+H]⁺

Anal. Calcd. for C₁₈H₁₅F₃N₂O₂S :

C, 56.84; H, 3.97; N, 7.36

Found. C, 56.44; H, 3.89; N, 7.19

30 Example 5

The following compounds described in (1)-(7) were prepared in a similar manner to that of Example 3.

(1) 1-(2-Fluorobenzyl)-5-[4-(methylsulfonyl)phenyl]-3-(trifluoromethyl)pyrazole
35

white crystals
mp : 136-138 °C
IR (KBr) : 1605, 1595, 1461 cm⁻¹
NMR (CDCl₃, δ) : 3.10 (3H, s), 5.43 (2H, s), 6.69
5 (1H, s), 6.90-7.40 (4H, m), 7.51 (2H, d, J=8Hz),
8.00 (2H, d, J=8Hz)
MS : 399 [M+H]⁺
Anal. Calcd. for C₁₈H₁₄F₄N₂O₂S :
C, 54.27; H, 3.54; N, 7.03
10 Found. C, 54.39; H, 3.52; N, 7.02.

(2) 1-(2,4-Difluorobenzyl)-5-[4-(methylsulfonyl)phenyl]-
3-(trifluoromethyl)pyrazole
white crystals
15 mp : 125-130 °C
IR (KBr) : 1612, 1508, 1467 cm⁻¹
NMR (CDCl₃, δ) : 3.10 (3H, s), 5.37 (2H, s), 6.67
10 (1H, s), 6.70-7.20 (3H, m), 7.52 (2H, d, J=8Hz),
8.02 (2H, d, J=8Hz)
20 MS : 417 [M+H]⁺
Anal. Calcd. for C₁₈H₁₃F₅N₂O₂S :
C, 51.93; H, 3.15; N, 6.73
Found. C, 51.68; H, 3.10; N, 6.64

25 (3) 1-(1-Naphthylmethyl)-5-[4-(methylsulfonyl)phenyl]-3-
(trifluoromethyl)pyrazole
white crystals
mp : 118-120 °C
IR (KBr) : 1600, 1506, 1465 cm⁻¹
30 NMR (CDCl₃, δ) : 3.01 (3H, s), 5.89 (2H, s), 6.73
(1H, s), 7.20-8.00 (11H, m)
MS : 431 [M+H]⁺
Anal. Calcd. for C₂₂H₁₇F₃N₂O₂S 1/3H₂O :
C, 60.54; H, 3.93; N, 6.42
35 Found. C, 60.77; H, 4.01; N, 6.39

(4) 4-[1-Benzyl-3-(trifluoromethyl)pyrazol-5-yl]benzene-sufonamide

white crystals

mp : 80-85 °C

5 IR (KBr) : 1714, 1563, 1463 cm^{-1}

NMR (CDCl_3 , δ) : 5.37 (2H, s), 6.66 (1H, s), 6.80-7.40 (5H, m), 7.43 (2H, d, $J=8\text{Hz}$), 7.97 (2H, d, $J=8\text{Hz}$)

MS : 382 $[\text{M}+\text{H}]^+$

10 Anal. Calcd. for $\text{C}_{17}\text{H}_{14}\text{F}_3\text{N}_3\text{O}_2\text{S}$:

C, 53.54; H, 3.70; N, 11.02

Found. C, 53.50; H, 3.90; N, 10.51

(5) 1-Benzyl-5-[3-fluoro-4-(methylsulfonyl)phenyl]-3-(trifluoromethyl)pyrazole

white crystals

mp : 118-120 °C

IR (KBr) : 1623, 1565, 1498, 1460 cm^{-1}

NMR (CDCl_3 , δ) : 3.25 (3H, s), 5.40 (2H, s), 6.69 (1H, s), 7.00-7.40 (7H, m), 7.99 (1H, m)

20 MS : 399 $[\text{M}+\text{H}]^+$

(6) 1-Benzyl-5-[4-(methylsulfonyl)phenyl]-3-(difluoromethyl)pyrazole

25 white crystals

mp : 105-108 °C

IR (KBr) : 1590, 1498, 1459 cm^{-1}

NMR (CDCl_3 , δ) : 3.08 (3H, s), 5.36 (2H, s), 6.65 (1H, s), 6.50-7.40 (6H, m), 7.50 (2H, d, $J=8\text{Hz}$),

30 7.98 (2H, d, $J=8\text{Hz}$)

MS : 363 $[\text{M}+\text{H}]^+$

(7) Ethyl 1-(2,4-difluorobenzyl)-5-[4-(methylthio)phenyl]pyrazole-3-carboxylate

35 pale yellow oil.

IR (neat) : 1742, 1610, 1504 cm^{-1}
NMR (CDCl_3 , δ) : 1.41 (3H, t, $J=7\text{Hz}$), 2.50 (3H, s),
4.43 (2H, q, $J=7\text{Hz}$), 6.60-7.30 (8H, m)
MS : 389 $[\text{M}+\text{H}]^+$

5

Example 6

A mixture of sodium methoxide (3.0 g), benzyl-hydrazine (3.67 g) and 3-[4-(methylthio)phenyl]-acrylonitrile (5.00 g) in methanol was stirred at 140°C for 1 hour. The reaction mixture was extracted with methylene chloride and the organic layer was washed with water and brine successively, dried, and concentrated to dryness. The obtained materials and manganese(VI) dioxide (MnO_2) (10.0 g) in ethyl acetate (200 ml) were refluxed for 2 hours. The reaction mixture was filtrated on celite and evaporated. The residue was chromatographed (toluene : ethyl acetate = 2 : 1) over silica gel to afford 1-benzyl-5-[4-(methylthio)phenyl]pyrazole-3-amine (0.98 g) as an orange crystals.

20 mp : 118-120 °C

IR (KBr) : 3432, 3351, 1602, 1548, 1504, 1459 cm^{-1}
NMR (DMSO-d_6 , δ) : 2.50 (3H, s), 5.23 (2H, s), 5.72 (1H, s), 6.90-7.40 (9H, m)
MS : 296 $[\text{M}+\text{H}]^+$

25

Example 7

A solution of sodium nitrite (890 mg) was added to an ice-salt-cooled mixture of 1-benzyl-5-[4-(methylthio)-phenyl]pyrazole-3-amine (2.50 g) in concentrated hydrochloric acid (25 ml). The resultant mixture was stirred at 0°C for 30 minutes and added dropwise to a mixture of copper(I) chloride (CuCl) (3.74 g) in concentrated hydrochloric acid (25 ml). The resultant mixture was stirred at room temperature and refluxed for 2 hours. To the reaction mixture was added a mixture of

ethyl acetate and water. The organic layer was washed with saturated sodium hydrogen carbonate, and brine successively, dried, and concentrated to dryness. The residue was chromatographed (toluene) over silica gel to afford 1-benzyl-3-chloro-5-[4-(methylthio)phenyl]pyrazole (0.46 g) as a red oil.

5 IR (neat) : 1600, 1550, 1482, 1444 cm^{-1}
NMR (CDCl_3 , δ) : 2.50 (3H, s), 5.25 (2H, s), 6.24 (1H, s), 6.90-7.40 (9H, m)

10 MS : 315 $[\text{M}+\text{H}]^+$

Example 8

A mixture of sodium methoxide (324 mg) and ethyl 1-(2,4-difluorobenzyl)-5-[4-(methylsulfonyl)phenyl]-pyrazole-3-carboxylate (840 mg) in formamide (5.0 ml) was warmed at 100°C for 1 hour. The reaction mixture was poured into ice-water and the precipitate was washed with water and ethanol to afford 1-(2,4-difluorobenzyl)-5-[4-(methylsulfonyl)phenyl]pyrazole-3-carboxamide (650 mg) as a white crystals.

20 mp : 205-208 °C
IR (KBr) : 3450, 3278, 3195, 1675, 1606, 1506, 1432 cm^{-1}
NMR (CDCl_3 , δ) : 3.10 (3H, s), 5.35 (2H, s), 5.5 (2H, bs), 6.70-7.10 (4H, m), 7.53 (2H, d, $J=8\text{Hz}$), 25 8.02 (2H, d, $J=8\text{Hz}$)
MS : 392 $[\text{M}+\text{H}]^+$

Example 9

30 A mixture of methanesulfonyl chloride (1.05 g) and 1-(2,4-difluorobenzyl)-5-[4-(methylsulfonyl)phenyl]-pyrazole-3-carboxamide (600 mg) in pyridine (5 ml) was warmed at 60°C for 2 hours. The reaction mixture was evaporated and ethyl acetate and water were added to the residue. The organic layer was washed with saturated

sodium hydrogen carbonate, and brine successively, dried, and concentrated to dryness. The residue was chromatographed (toluene : ethyl acetate = 5 : 1) over silica gel to afford 1-(2,4-difluorobenzyl)-5-[4-(methylsulfonyl)phenyl]pyrazole-3-carbonitrile (263 mg) as a white crystals.

mp : 103-104 °C
IR (KBr) : 2242, 1610, 1508, 1428 cm⁻¹
NMR (CDCl₃, δ) : 3.11 (3H, s), 5.35 (2H, s), 6.70-7.20 (4H, m), 7.53 (2H, d, J=8Hz), 8.05 (2H, d, J=8Hz)
MS : 374 [M+H]⁺

Example 10

Sodium hydride (60%, 338 mg) was added to a solution of 4-(4-fluorophenyl)-3-[4-(methylthio)phenyl]pyrazole (1.6 g) in dimethylformamide (30 ml) at 0°C. The resultant mixture was stirred at same temperature for 1 hour and benzylchloride (1.06 g) was dropwise to the above mixture. The reaction mixture was stirred for additional 2 hours at ambient temperature and poured into ice-water and extracted with ethyl acetate. The organic layer was washed with water, and brine successively, dried, and concentrated to dryness. The residue was chromatographed (CH₂Cl₂ : n-hexane = 3 : 2) over silica gel to afford 1-benzyl-4-(4-fluorophenyl)-5-[4-(methylthio)phenyl]pyrazole (156 mg) from second fraction.

IR (nujol) : 1550, 1530, 1250 cm⁻¹
NMR (DMSO-d₆, δ) : 2.50 (3H, s), 5.02 (2H, s), 6.95-7.33 (13H, m), 7.89 (1H, s)
MS : 375 [M+H]⁺

Example 11

The following compounds described in (1)-(3) were prepared in a similar manner to that of Example 10.

(1) 1-(4-Chlorobenzyl)-4-(4-fluorophenyl)-5-[4-(methylthio)phenyl]pyrazole

IR (nujol) : 1600, 1550 cm^{-1}

NMR (DMSO-d₆, δ) : 2.50 (3H, s), 5.19 (2H, s), 6.70-7.43 (12H, m), 7.91 (1H, s)

5

(2) 4-(4-Fluorophenyl)-1-(4-nitrobenzyl)-5-[4-(methylthio)phenyl]pyrazole

IR (nujol) : 1600, 1560, 1510 cm^{-1}

NMR (DMSO-d₆, δ) : 2.48 (3H, s), 5.35 (2H, s), 7.00-7.40 (10H, m), 7.96 (1H, s), 8.16 (2H, d, $J=8\text{Hz}$)
MS : 419 [M+H]⁺

10

(3) 4-(4-Fluorophenyl)-1-(4-methoxybenzyl)-5-[4-(methylthio)phenyl]pyrazole

15

IR (nujol) : 1600, 1580 cm^{-1}

NMR (DMSO-d₆, δ) : 3.33 (3H, s), 3.79 (3H, s), 5.11 (2H, s), 6.70-7.40 (12H, m), 7.86 (1H, s)

MS : 405 [M+H]⁺

20

Example 12

The following compounds described in (1)-(6) were prepared in a similar manner to that of Example 2.

25 (1) Ethyl 1-(2,4-difluorobenzyl)-5-[4-(methylsulfonyl)phenyl]pyrazole-3-carboxylate

white crystals

mp : 134-137 $^{\circ}\text{C}$

IR (KBr) : 1708, 1612, 1509, 1463 cm^{-1}

30 NMR (CDCl₃, δ) : 1.42 (3H, t, $J=7\text{Hz}$), 3.09 (3H, s), 4.45 (2H, q, $J=7\text{Hz}$), 6.70-7.20 (4H, m), 7.50 (2H, d, $J=8\text{Hz}$), 8.00 (2H, d, $J=8\text{Hz}$)

MS : 421 [M+H]⁺

35 (2) 1-Benzyl-3-chloro-5-[4-(methylsulfonyl)phenyl]-

pyrazole

white crystals

mp : 135-138 °C

5 IR (KBr) : 1542, 1533, 1482, 1448 cm⁻¹
NMR (CDCl₃, δ) : 3.08 (3H, s), 5.27 (2H, s), 6.36
(1H, s), 7.00-7.40 (5H, m), 7.49 (2H, d, J=8Hz),
7.97 (2H, d, J=8Hz)
MS : 347 (M⁺¹), 349 (M⁺³)

10 (3) 1-Benzyl-4-(4-fluorophenyl)-5-[4-(methylsulfonyl)-phenyl]pyrazole

mp : 173-174 °C

15 IR (nujol) : 1550, 1530, 1310 cm⁻¹
NMR (DMSO-d₆, δ) : 3.28 (3H, s), 5.25 (2H, s),
6.94-7.28 (9H, m), 7.54 (2H, d, J=8Hz), 7.93
(1H, s), 7.98 (2H, d, J=8Hz).
MS : 407 [M+H]⁺

20 (4) 1-(4-Chlorobenzyl)-4-(4-fluorophenyl)-5-[4-(methylsulfonyl)phenyl]pyrazole

mp : 166-167 °C

25 IR (nujol) : 1605, 1550 cm⁻¹
NMR (DMSO-d₆, δ) : 3.28 (3H, s), 5.25 (2H, s), 6.98-7.55 (10H, m), 7.91 (2H, d, J=8Hz), 7.95 (1H, s)
MS : 441 [M+H]⁺

(5) 4-(4-Fluorophenyl)-1-(4-nitrobenzyl)-5-[4-(methylsulfonyl)phenyl]pyrazole

mp : 220-221 °C

30 IR (nujol) : 1600, 1510, 1340 cm⁻¹
NMR (DMSO-d₆, δ) : 3.32 (3H, s), 5.40 (2H, s), 7.00-7.20 (6H, m), 7.22 (2H, d, J=8Hz), 7.98 (2H, d, J=8Hz), 8.00 (1H, s), 8.14 (2H, d, J=8Hz)
MS : 452 [M+H]⁺

(6) 4-(4-Fluorophenyl)-1-(4-methoxybenzyl)-5-[4-(methylsulfonyl)phenyl]pyrazole

mp : 167-168 °C

IR (nujol) : 1600, 1550, 1510 cm^{-1}

5 NMR (DMSO-d₆, δ) : 3.29 (3H, s), 3.69 (3H, s), 5.17 (2H, s), 6.80-7.20 (8H, m), 7.54 (2H, d, J=8Hz), 7.90 (1H, s), 7.99 (2H, d, J=8Hz)

MS : 437 [M+H]⁺

10 Example 13

N-Chlorosuccinimide (25 mg) and 1-benzyl-4-(4-fluorophenyl)-5-[4-(methylsulfonyl)phenyl]pyrazole (50 mg) in acetic acid (5 ml) was stirred at 70°C for 1 hour. The reaction mixture was stirred for additional 30 minutes at 15 75°C and poured into ice-water and extracted with ethyl acetate. The organic layer was washed with water, and brine successively, dried, and concentrated to dryness. The residue was recrystallized from ethanol to afford 20 1-benzyl-3-chloro-4-(4-fluorophenyl)-5-[4-(methylsulfonyl)phenyl]pyrazole (35 mg).

mp : 156-157 °C

IR (nujol) : 1600, 1550, 1510, 1300 cm^{-1}

25 NMR (DMSO-d₆, δ) : 3.26 (3H, s), 5.25 (2H, s), 7.01-7.34 (9H, m), 7.56 (2H, d, J=8Hz), 7.95 (2H, d, J=8Hz)

MS : 441 [M+H]⁺

Example 14

30 The following compounds described in (1)-(2) were prepared in a similar manner to that of Example 13.

(1) 3-Chloro-1-(4-chlorobenzyl)-4-(4-fluorophenyl)-5-[4-(methylsulfonyl)phenyl]pyrazole.

mp : 148-150 °C

35 IR (nujol) : 1600 cm^{-1}

NMR (DMSO-d₆, δ) : 3.26 (3H, s), 5.25 (2H, s), 7.00-
7.60 (10H, m), 7.91 (2H, d, J=8Hz)

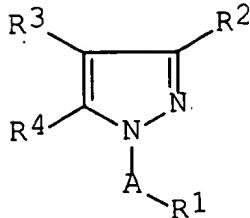
MS : 476 [M+H]⁺

5 (2) 3-Chloro-4-(4-fluorophenyl)-1-(4-nitrobenzyl)-5-
[4-(methylsulfonyl)phenyl]pyrazole
mp : 130-131 °C
IR (nujol) : 1600, 1570, 1550 cm⁻¹
NMR (DMSO-d₆, δ) : 3.25 (3H, s), 5.42 (2H, s), 7.00-
7.30 (6H, m), 7.56 (2H, d, J=8Hz), 7.95 (2H, d,
J=8Hz), 8.17 (2H, d, J=8Hz)
MS : 486 [M+H]⁺

C L A I M S

1. A compound of the formula :

5



wherein R¹ is aryl optionally substituted with
10 substituent(s) selected from the group
consisting of halogen, nitro, amino,
esterified carboxy, carboxy, carbamoyl,
cyano and lower alkoxy,

R² is hydrogen, amino, halogen, esterified
15 carboxy, carboxy, carbamoyl, cyano, or
lower alkyl substituted with halogen,

R³ is hydrogen, aryl optionally substituted
20 with halogen, or lower alkyl optionally
substituted with hydroxy, amino or
carboxy,

R⁴ is aryl substituted with substituent(s)
selected from the group consisting of
halogen, lower alkoxy, lower alkylthio,
lower alkylsulfinyl, lower
25 alkylsulfonyl, sulfamoyl, and lower
alkylsulfamoyl, and

A is lower alkylene,
and its salt.

30 2. The compound according to claim 1,
wherein A is methylene.

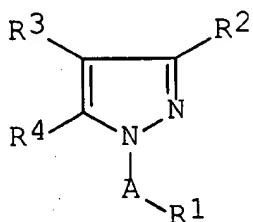
3. The compound according to claim 2,
35 wherein R² is amino, halogen, esterified carboxy,
carboxy, carbamoyl, cyano, or lower

alkyl substituted with halogen, and
R³ is hydrogen.

4. The compound according to claim 2,
5 wherein R² is hydrogen or halogen, and
R³ is aryl optionally substituted with
halogen, or lower alkyl optionally
substituted with hydroxy, amino or
carboxy.

10

5. A process for preparing a compound of the formula :



15

wherein R¹ is aryl optionally substituted with
substituent(s) selected from the group
consisting of halogen, nitro, amino,
esterified carboxy, carboxy, carbamoyl,
20 cyano and lower alkoxy,

20

R² is hydrogen, amino, halogen, esterified
carboxy, carboxy, carbamoyl, cyano, or
lower alkyl substituted with halogen,

25

R³ is hydrogen, aryl optionally substituted
with halogen, or lower alkyl optionally
substituted with hydroxy, amino or
carboxy,

30

R⁴ is aryl substituted with substituent(s)
selected from the group consisting of
halogen, lower alkoxy, lower alkylthio,
lower alkylsulfinyl, lower
alkylsulfonyl, sulfamoyl, and lower
alkylsulfamoyl, and

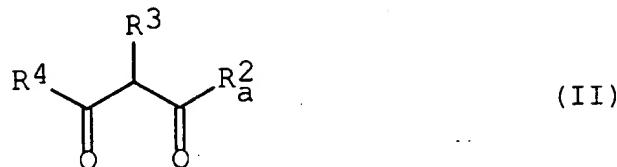
35

A is lower alkylene,

or its salt, which comprises,

(1) reacting a compound of the formula :

5

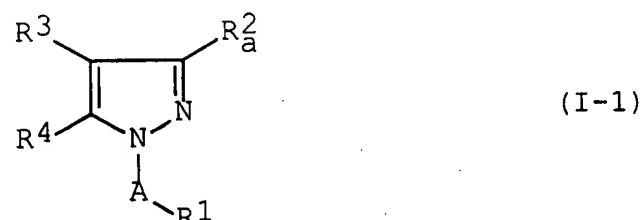


10



15

or its salt, to give a compound of the formula :



20

or its salt, in the above formulas

R^1 , R^3 , R^4 and A are each as defined above, and
 R^2_a is hydrogen, esterified carboxyl, carbamoyl,
cyano, or lower alkyl substituted with
halogen,

25

(2) reacting a compound of the formula :



30

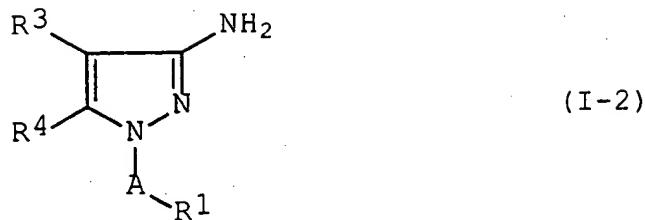
or its salt, with a compound of the formula :



35

or its salt, to give a compound of the formula :

5

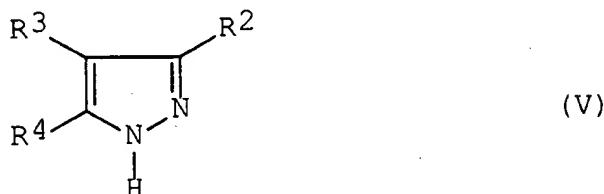


or its salt, in the above formulas

R^1 , R^3 , R^4 and A are each as defined above,

10 (3) reacting a compound of the formula :

15



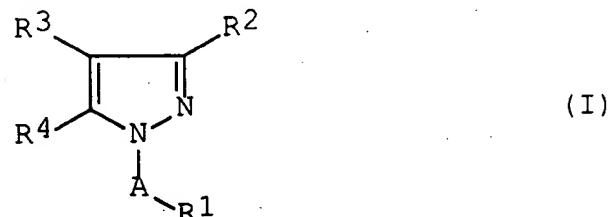
or its salt, with a compound of the formula :

20



or its salt, to give a compound of the formula :

25



or its salt, in the above formulas

R^1 , R^2 , R^3 , R^4 and A are each as defined above,

30

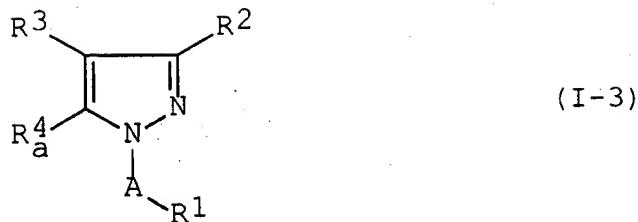
and

X^1 is leaving group,

(4) oxidizing a compound of the formula :

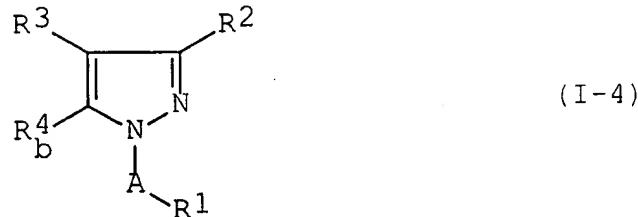
35

5



or its salt, to give a compound of the formula :

10



or its salt, in the above formulas

15

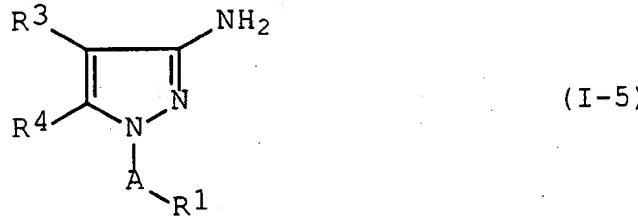
R^1 , R^2 , R^3 and A are each as defined above,

R^4_a is aryl substituted with lower alkylthio, and

R^4_b is aryl substituted with lower alkylsulfinyl
or lower alkylsulfonyl,

(5) subjecting a compound of the formula :

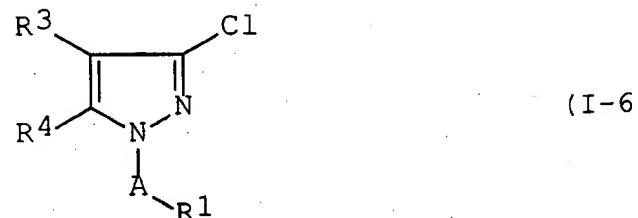
20



25

or its salt, to chlorination to give a compound of the formula :

30

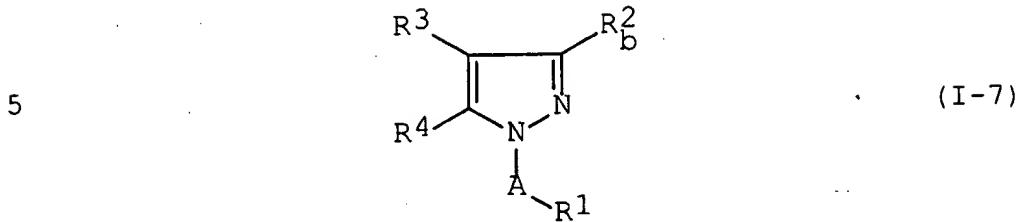


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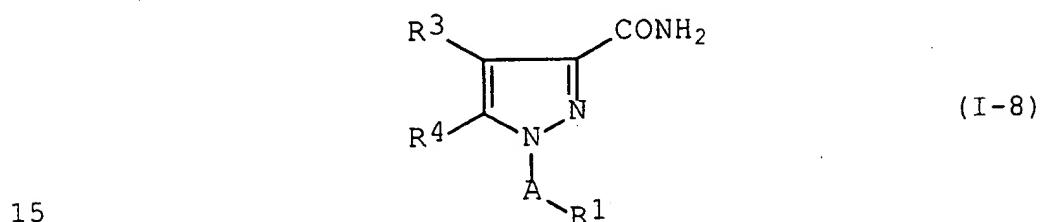
or its salt, in the above formulas

R^1 , R^2 , R^3 and A are as defined above,

(6) subjecting a compound of the formula :



10 or its salt, to amidation reaction, to give a compound
of the formula :



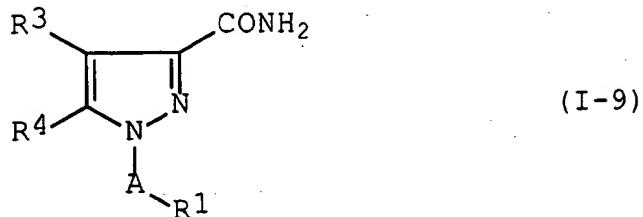
or its salt, in the above formulas

R^1 , R^3 , R^4 and A are each as defined above, and
 R^2_b is esterified carboxy or carboxy, or

20

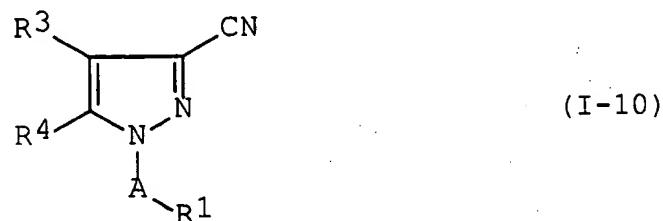
(7) subjecting a compound of the formula :

25



or its salt, to dehydration reaction, to give a
compound of the formula :

30



35

or its salt, in the above formulas
R¹, R³, R⁴ and A are each as defined above.

6. A pharmaceutical composition comprising the compound
5 of claim 1, as an active ingredient, in association
with a pharmaceutically non-toxic carrier or
excipient.
7. A compound of claim 1 for use as a medicament.
10
8. COX-II inhibiting agent comprising the compound of
claim 1.
9. A method for the treatment and/or prevention of
15 inflammatory conditions, various pains, collagen
diseases, autoimmune diseases, various immunity
diseases, analgesic, thrombosis, cancer or
neurodegenerative diseases which comprises administering
an effective amount of the compound of claim 1 to
20 human beings or animals.
10. Use of the compound of claim 1 for the manufacture of
a medicament for treatment and/or prevention of
inflammatory conditions, various pains, collagen
25 diseases, autoimmune diseases, various immunity
diseases, analgesic, thrombosis, cancer or
neurodegenerative diseases in human beings or animals.

INTERNATIONAL SEARCH REPORT

International Application No
PCT/JP 98/05041

A. CLASSIFICATION OF SUBJECT MATTER
IPC 6 C07D231/12 A61K31/415

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
IPC 6 C07D

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	WO 97 13755 A (FUJISAWA PHARMACEUTICAL CO., LTD) 17 April 1997 see page 1 - page 2 ---	1,6
Y	WO 95 15318 A (G.D SEARLE &CO.) 8 June 1995 cited in the application see page 4 - page 29 ---	1,6
Y	WO 95 15316 A (G. D. SEARLE & CO.) 8 June 1995 cited in the application see page 4 - page 53, line 28 ---	1,6
Y	EP 0 418 845 A (FUJISAWA PHARMACEUTICAL CO. LTD.) 27 March 1991 cited in the application see page 2 - page 3, line 10 ---	1,6
-/-		

Further documents are listed in the continuation of box C.

Patent family members are listed in annex.

* Special categories of cited documents :

- "A" document defining the general state of the art which is not considered to be of particular relevance
- "E" earlier document but published on or after the international filing date
- "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- "O" document referring to an oral disclosure, use, exhibition or other means
- "P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

"&" document member of the same patent family

Date of the actual completion of the international search

30 March 1999

Date of mailing of the international search report

08/04/1999

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Fax: (+31-70) 340-3016

Authorized officer

Kyriakakou, G

INTERNATIONAL SEARCH REPORTInternational Application No
PCT/JP 98/05041**C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT**

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	DE 41 26 543 A (CHEMISCHE UND PHARMAZEUTISCHE FABRIKEN FAHLBERG-LIST GMBH) 11 February 1993 see the whole document -----	1,6
Y	US 5 486 534 A (LEN F. LEE ET AL.) 23 January 1996 see the whole document -----	1,6

INTERNATIONAL SEARCH REPORT

International application No.

PCT/JP 98/05041

Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. Claims Nos.: because they relate to subject matter not required to be searched by this Authority, namely:

Although claim 9 is directed to a method of treatment of the human/animal body, the search has been carried out and based on the alleged effects of the compound/composition.
2. Claims Nos.: because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:
3. Claims Nos.: because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.
2. As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:
4. No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims: it is covered by claims Nos.:

Remark on Protest

The additional search fees were accompanied by the applicant's protest.

No protest accompanied the payment of additional search fees.

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/JP 98/05041

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INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/JP 98/05041

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